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(54) **POROUS FLAME HOLDER FOR LOW NOX COMBUSTION**

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CPC **F23D 14/66** (2013.01); **F23C 3/002** (2013.01); **F23D 14/14** (2013.01); **F23D 14/64** (2013.01);

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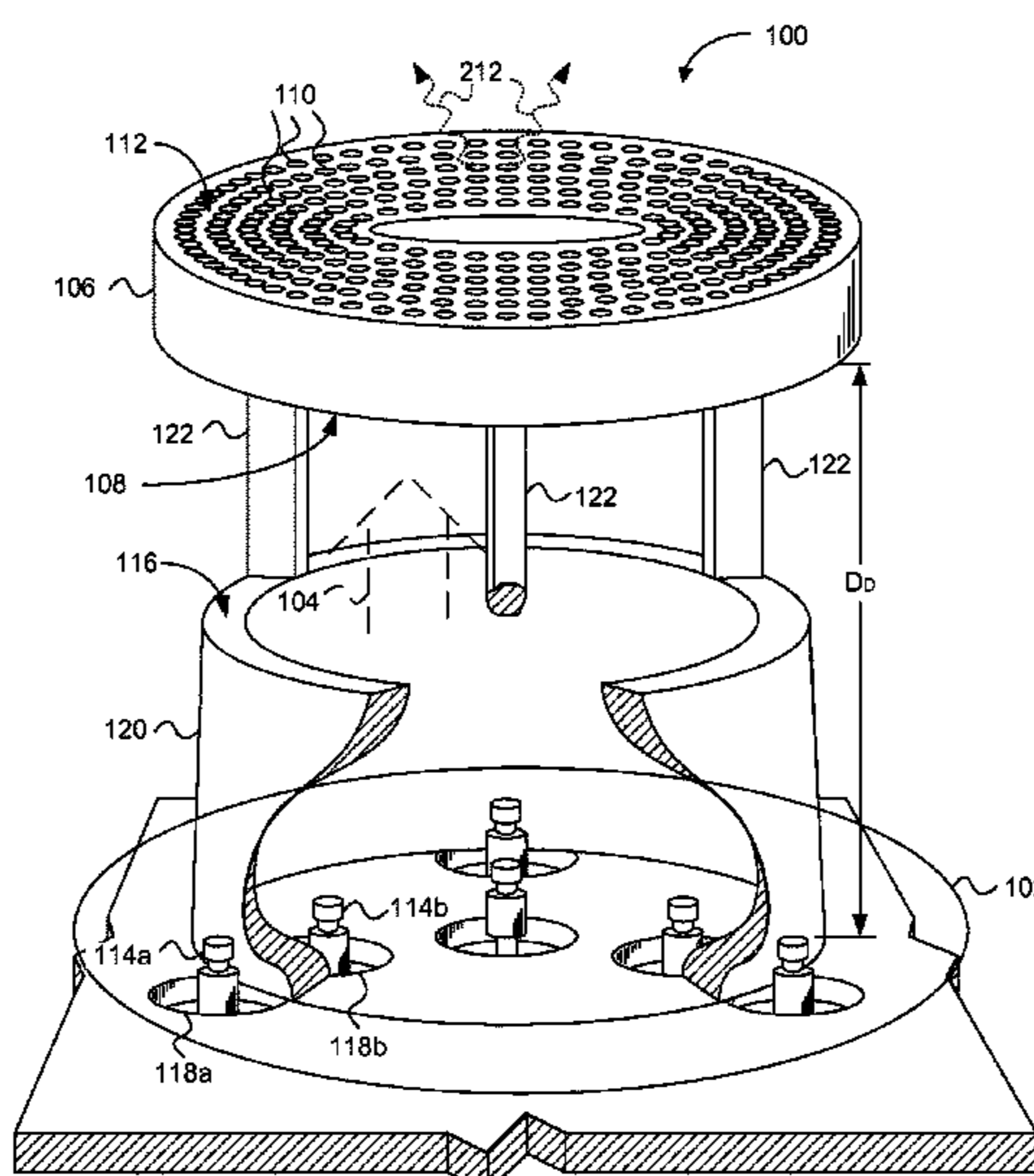
(57) **ABSTRACT**

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A burner includes a porous flame holder configured to support a combustion reaction to achieve a very low output of oxides of nitrogen (NOx).

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30 Claims, 6 Drawing Sheets



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FIG. 1A

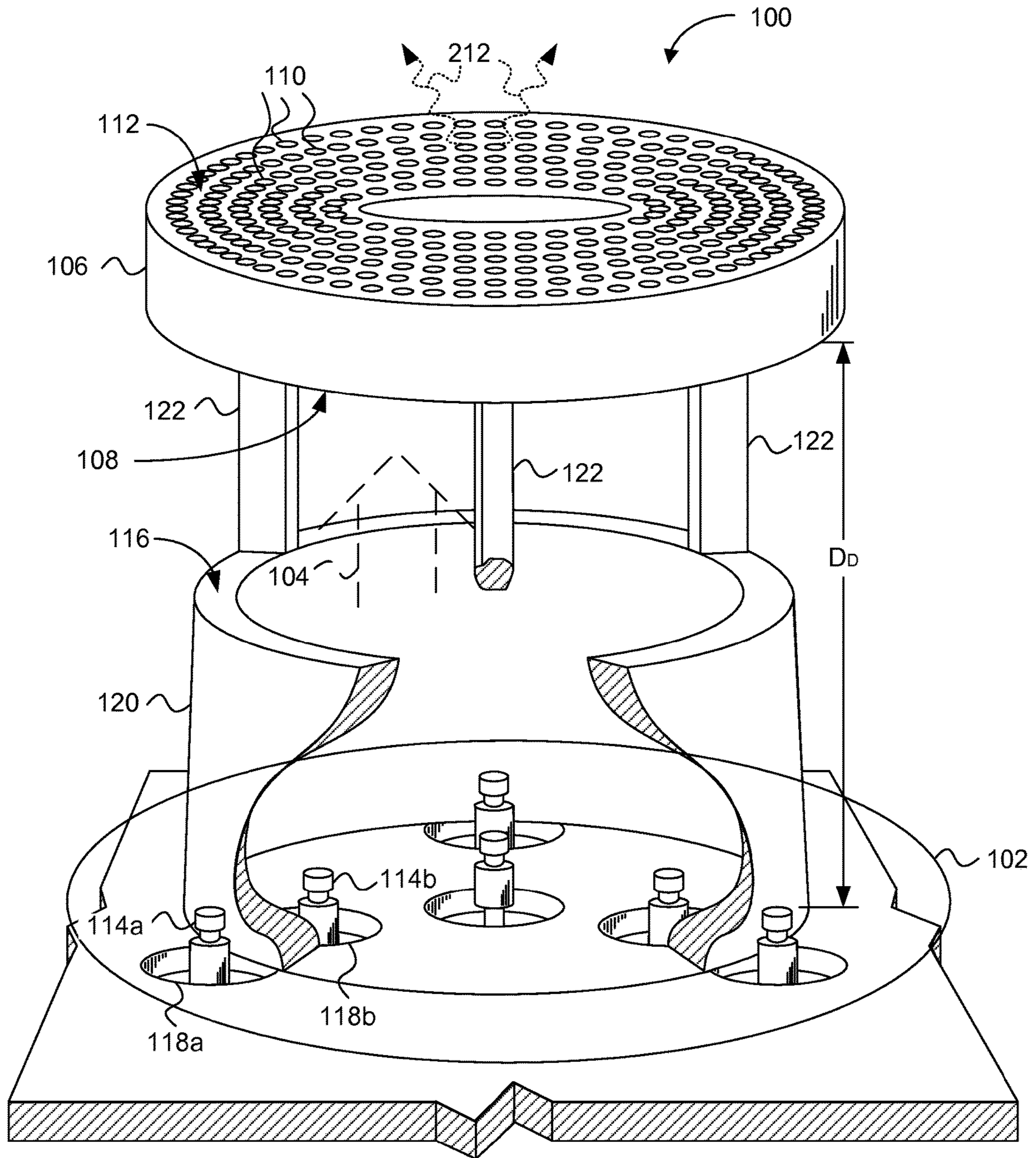


FIG. 1B

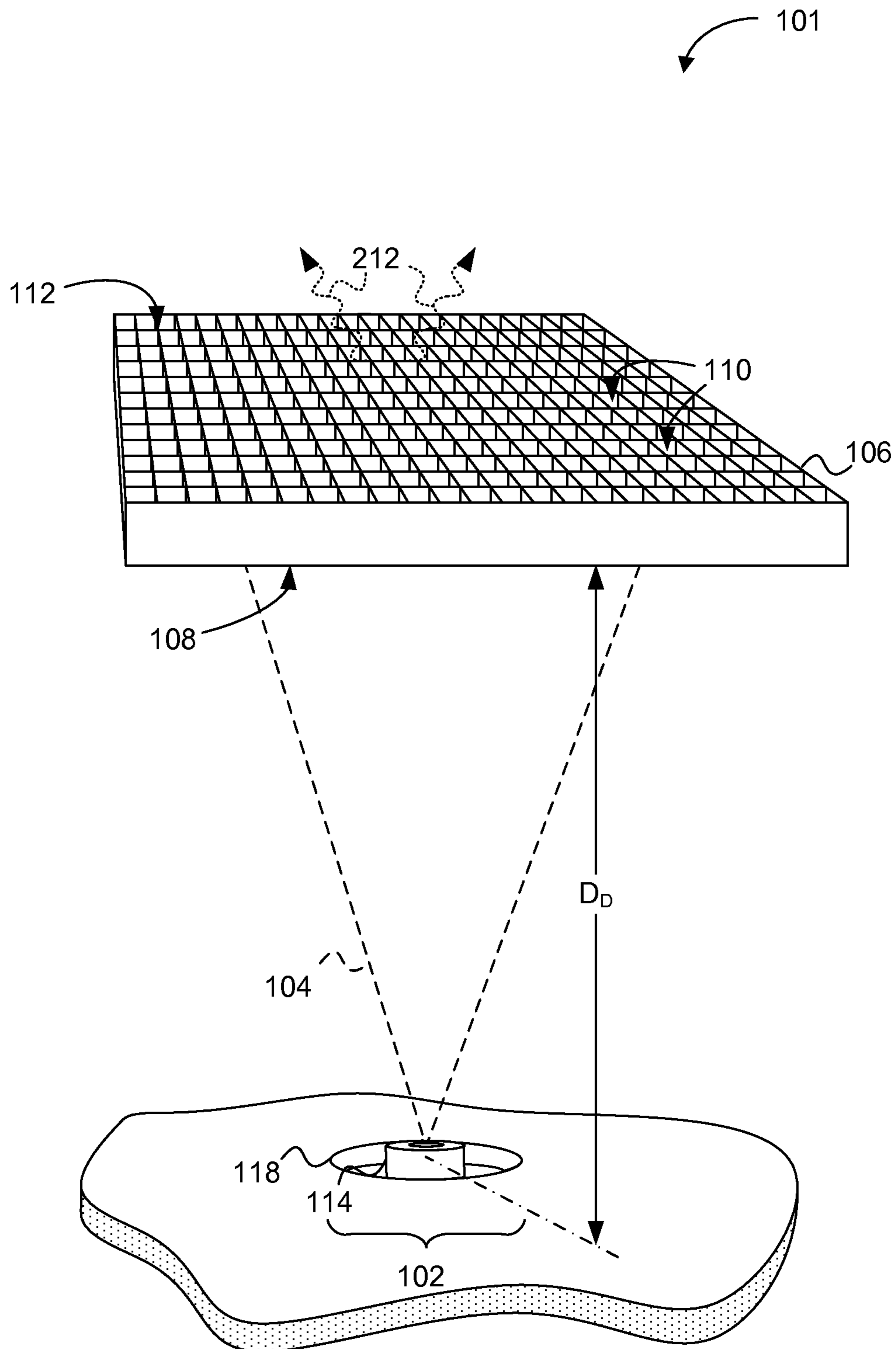


FIG. 2

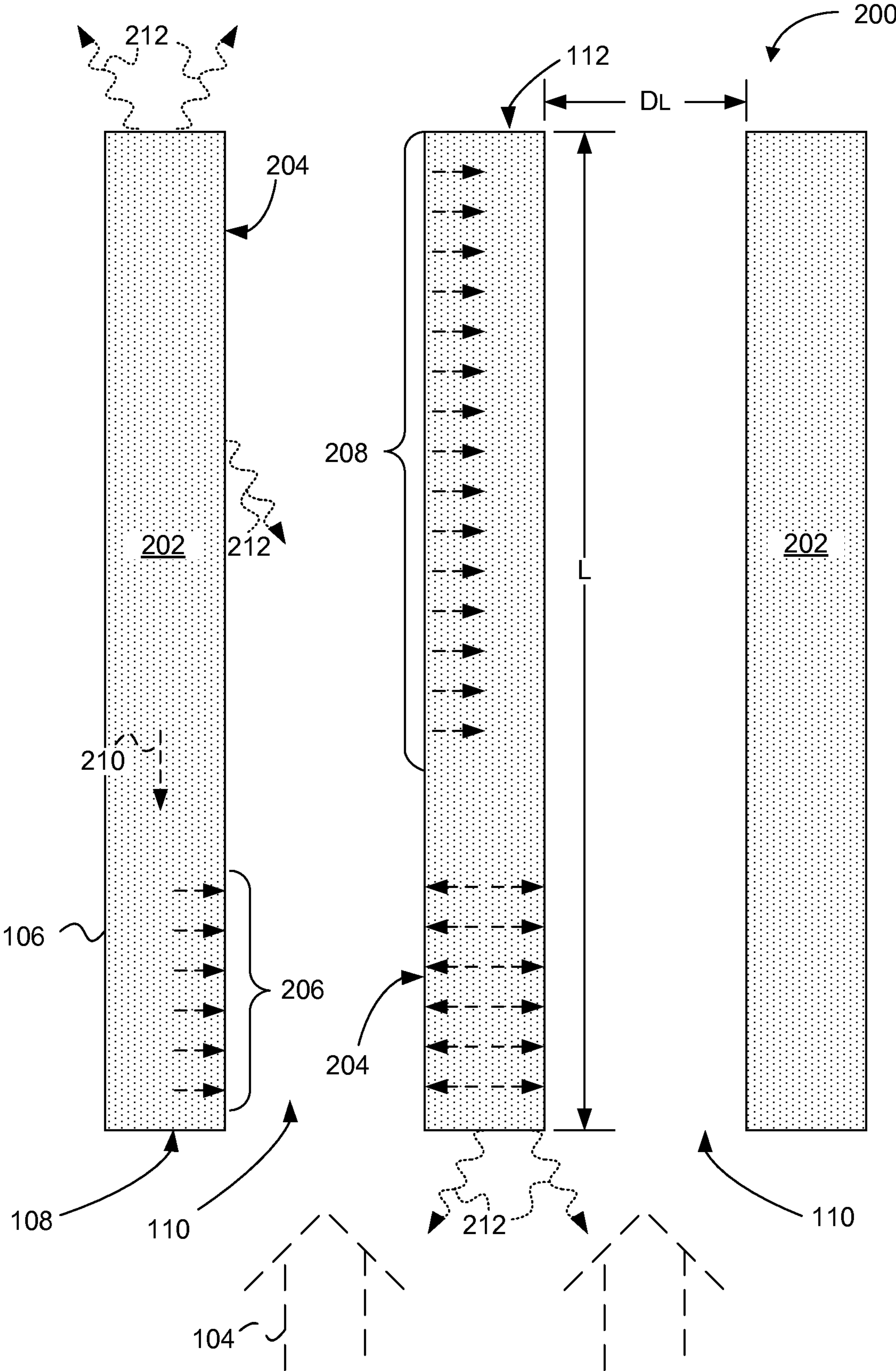


FIG. 3

300

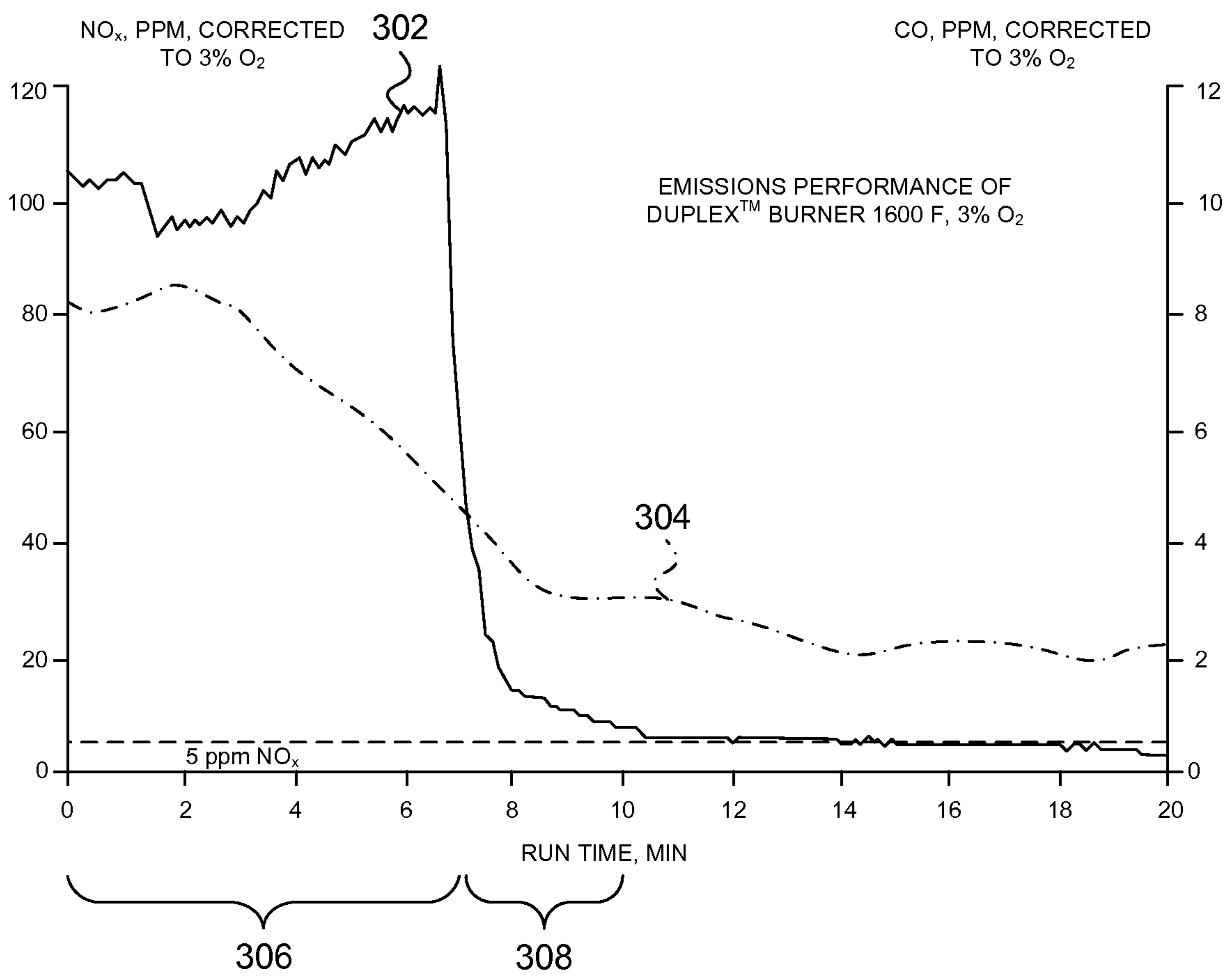


FIG. 4

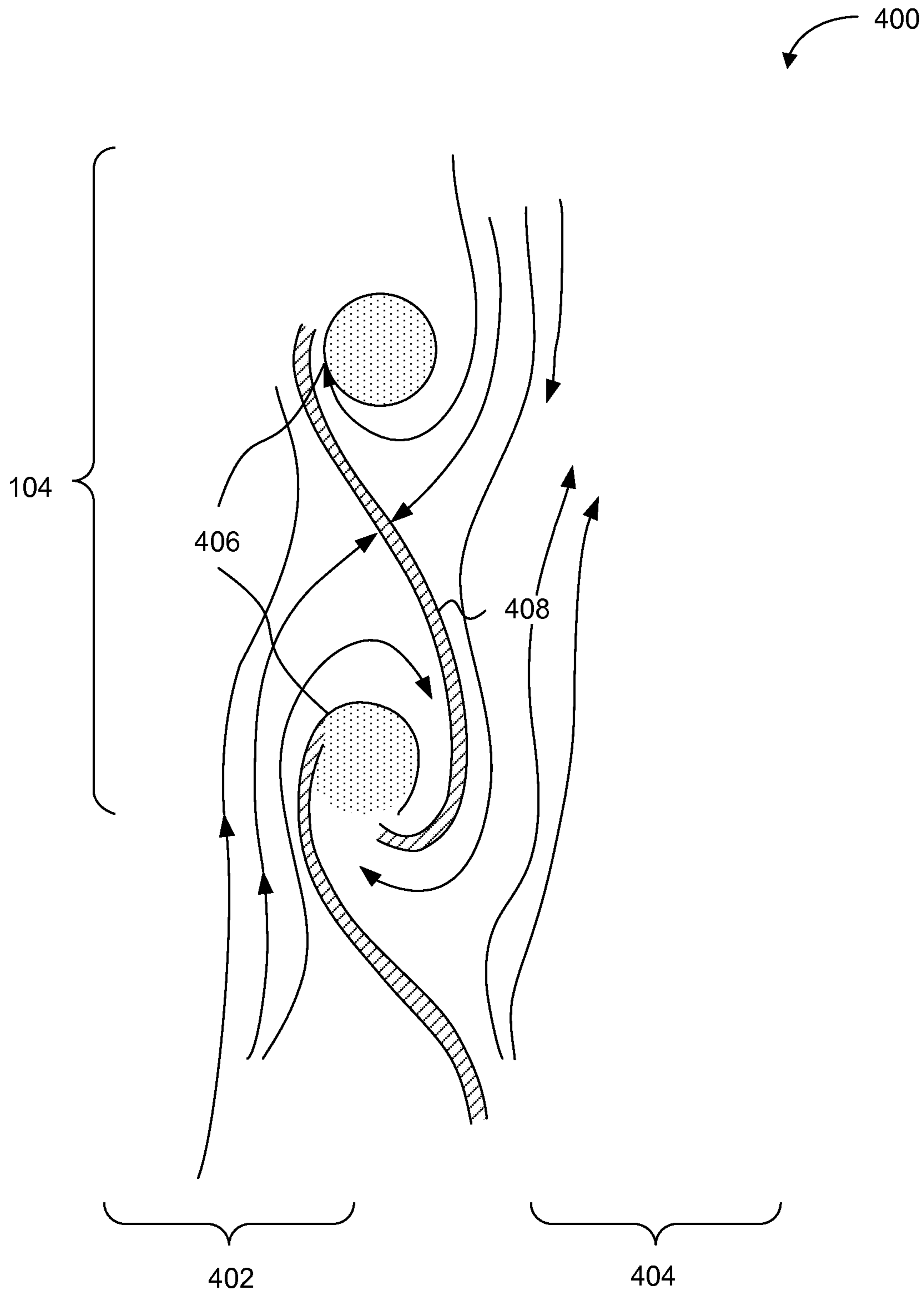
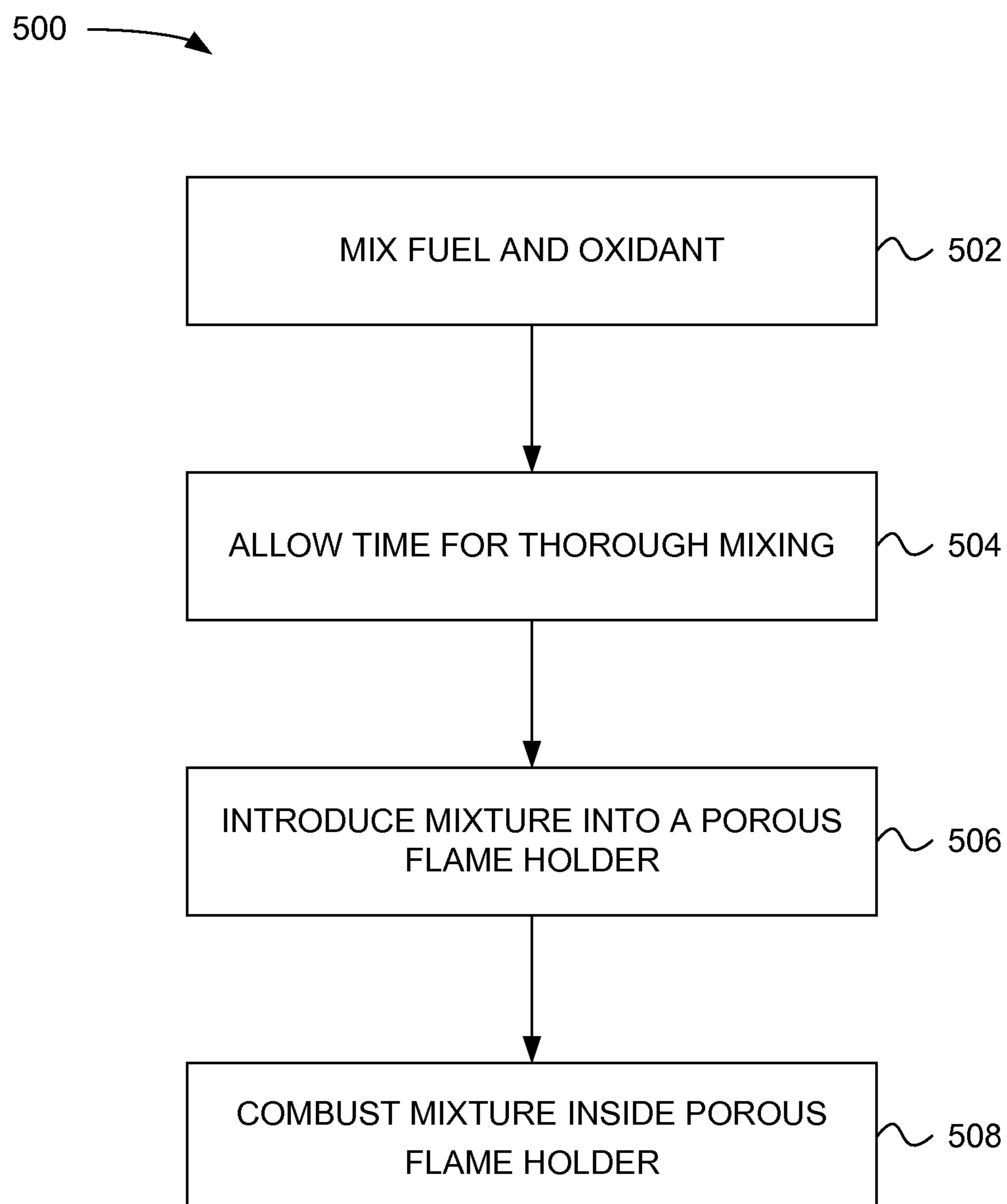


FIG. 5



POROUS FLAME HOLDER FOR LOW NOX COMBUSTION

CROSS-REFERENCE TO RELATED APPLICATIONS

The present application is a U.S. National Phase application under 35 U.S.C. 371 of co-pending International Patent Application No. PCT/US2014/057072, entitled "POROUS FLAME HOLDER FOR LOW NOx COMBUSTION", filed Sep. 23, 2014; which application claims the priority benefit of U.S. Provisional Patent Application No. 61/887,741, entitled "POROUS FLAME HOLDER FOR LOW NOx COMBUSTION", filed Oct. 7, 2013; and U.S. Provisional Patent Application No. 61/881,368, entitled "PROGRESS AND RECENT ADVANCES USING ELECTRODYNAMIC COMBUSTION CONTROL (ECC)", filed Sep. 23, 2013; which is a Continuation-in-part of International Patent Application No. PCT/US2014/016632, entitled "FUEL COMBUSTION SYSTEM WITH A PERFORATED REACTION HOLDER", filed Feb. 14, 2014; each of which, to the extent not inconsistent with the disclosure herein, is incorporated herein by reference.

BACKGROUND

NOx is a pollutant regulated by the EPA and a key metric of burner performance. Strict new NOx control regulations are being implemented in several regions of the country including Texas and California. California's South Coast Air Quality Management District's Rule 1146 required that burners produce less than 9 ppm of NOx no later than July 2014. Industry groups anticipate that these limits will soon be further reduced to as low as 5 ppm in some areas with the rest of the country to follow suit.

To address this challenge, some burner and combustion system manufacturers have been able to develop systems that can achieve the NOx targets, but inherent design tradeoffs impose high costs to energy efficiency that become prohibitive at these very low emissions levels, even with natural gas at historically low prices.

The biggest cost associated with prior art Low- and Ultra-Low NOx burners has been the significant loss in energy efficiency that results. This loss stems directly from the combined effect of recirculating flue gas and increasing excess air to cool the combustion reaction along with a loss of turn-down because of flame instability, and can result in substantial increases in fuel consumption and parasitic power losses of up to 20-30%.

The market has long preferred low NOx and Ultra-Low NOx burners to more costly post-combustion treatment alternatives such as Selective Catalytic Reduction (SCR) systems that are more costly to install, complex to operate and consume considerable quantities of hazardous materials such as anhydrous ammonia.

SUMMARY

An embodiment demonstrated a 95% reduction in NOx emissions (down to less than 2 ppm) using a prototype burner based on a porous flame holder in a furnace operating at a temperature of ~1600 F with O₂ concentrations ranging from 2.5% to 3.2% and virtually no CO.

Embodiments achieve reductions in NOx to 2 ppm without costly FGR or SCR systems. In addition, flame length is reduced by up to 80%. And, unlike other low NOx systems, the burner keeps oxygen at normal operating levels (1-3%),

and can maintain a stable combustion reaction throughout a wide operating range. Because of these features, the burner enables significant advantages in fuel efficiency and process throughput as compared to conventional Low- and Ultra-low-NOx burners.

Many traditional Low-NOx burners make use of increased flame length and reduced momentum to reduce NOx, but the resulting poor 'flame pattern' can cause combustion reaction impingement and coking. Refinery process heaters are particularly sensitive to this problem, due to the direct negative impact on product throughput and plant revenue as the firing rate must be throttled to avoid equipment damage. According to our analysis, a 3% to 7% loss in firing capacity due to poor flame pattern can cost millions of dollars annually in lost process throughput. Removing this bottleneck could improve plant profitability by between \$12 and \$28 million per plant, per year. This is of particularly high value because it leverages so much capital plant by increasing capacity.

According to an embodiment, a burner includes a fuel nozzle assembly, configured to output a gaseous mixture including fuel and oxidant, and a porous flame holder defining a plurality of gas passages, each having a minimum dimension equal to or greater than a fuel quenching distance, aligned to receive the gaseous mixture, the gas passages being configured to pass a combustion reaction supported by the gaseous mixture. The gas passages each include a wall configured to receive heat from a reacting portion of the gaseous mixture, radiate and/or conduct the heat toward an unreacted portion of the gaseous mixture, and output the heat adjacent to the unreacted portion of the gaseous mixture to heat the gaseous mixture. The porous flame holder maintains stable combustion within the gas passages. The heat provided to the mixture in the gas passages can keep the combustion reaction stable even at fuel/oxidant mixtures at or below a lean flammability limit that would be stable in a conventional burner.

According to an embodiment, a method of lowering combustion NOx includes outputting fuel and oxidant to a burner, allowing time for the mixture to evolve to a better-mixed state, and then combusting the mixture inside a porous flame holder. The porous flame holder is configured to receive heat from the combustion reaction in each of a plurality of gas passages inside the porous flame holder, and conduct heat to the mixture sufficiently to cause stable combustion to be supported inside the porous flame holder.

According to an embodiment, a diluted fuel burner includes a fuel nozzle assembly configured to output a gaseous mixture including fuel and oxidant, a porous flame holder defining gas passages aligned to receive the gaseous mixture, the gas passages being configured to carry a combustion reaction supported by the gaseous mixture, and a porous flame holder support structure configured to hold the porous flame holder away from the fuel nozzle assembly at a dilution distance selected to allow dilution of the gaseous mixture.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1A is a cutaway view of a burner, according to an embodiment.

FIG. 1B is a diagram of a burner, according to another embodiment.

FIG. 2 is a side sectional diagram of a plurality of gas passages included in the porous flame holder portion of the burner of FIG. 1, according to an embodiment.

FIG. 3 is a graph of the performance of burner NO_x and CO emissions performance, according to an embodiment.

FIG. 4 is a diagram of fuel and combustion air mixing used to produce a gaseous mixture, according to an embodiment.

FIG. 5 is a flow chart showing a method for using the apparatus shown in FIGS. 1A, 1B and 2, according to an embodiment.

DETAILED DESCRIPTION

In the following detailed description, reference is made to the accompanying drawings, which form a part hereof. In the drawings, similar symbols typically identify similar components, unless context dictates otherwise. The illustrative embodiments described in the detailed description, drawings, and claims are not meant to be limiting. Other embodiments may be utilized, and other changes may be made, without departing from the spirit or scope of the subject matter presented here.

FIG. 1A is a diagram of a burner 100, according to an embodiment. FIG. 1B is a diagram of a burner 101, according to another embodiment. Referring to FIGS. 1A and 1B, The burner 100, 101 includes a fuel nozzle assembly 102 configured to output a gaseous mixture 104 including fuel and oxidant. A porous flame holder 106 is aligned to receive the gaseous mixture 104 at an upstream surface 108. The porous flame holder 106 defines a plurality of gas passages 110, each of the gas passages 110 being configured to carry a combustion reaction supported by the gaseous mixture 104. In some embodiments, each of the gas passages 110 can have a minimum lateral dimension equal to or greater than a flame quenching distance.

FIG. 2 is a side sectional diagram 200 of a plurality of gas passages 110 included in the porous flame holder 106 portion of the burners 100, 101 of FIGS. 1A and 1B, according to an embodiment. A minimum lateral dimension D_L is illustrated. Flame quenching distance refers to a lateral dimension through which a flame will not pass. There are two senses in which the gas passages 106 have dimensions D greater than the quenching distance. In both senses, the flame is kept in a state of ignition within each gas passage and is allowed to pass freely therethrough (although thermal and aerodynamic effects described below typically confine the combustion reaction to occur substantially within the gas passages 110, between the upstream surface 108 and a downstream surface 112 of the porous flame holder 106).

In the classical sense, the quenching distance is a tabulated value that is determined under stoichiometric conditions. It is generally considered a property of the fuel and exists as a tabulated property. Most hydrocarbons have quenching distances of about 0.1". For example, NACA Lewis Report 1300 tabulates quenching distance as shown in Table 1.

TABLE 1

FUEL QUENCHING DISTANCES	
HYDROCARBON FUEL	QUENCHING DISTANCE
n-Butane	0.12"
Methane	0.10"
Propane	0.08"
Hydrogen	0.025"

The quenching distance represents the diameter of an orifice such that a stoichiometrically premixed flame cannot

propagate upstream through the orifice into a premix reservoir. The mechanism is essentially one of heat abstraction—the flame giving up too much energy as it attempts to flashback through the orifice.

In contrast to porous flame holders 106 described herein, radiant burners that support surface combustion must have a minimum pore size less than the quenching distance for the particular fuel and temperature to avoid flashback, and it could be considered a tautology that if the flame flashes back, the pore size must be greater than the actual quenching distance under the operating conditions.

In a second sense, the porous flame holder 106 includes gas passages 110 that are larger than an operating quenching distance by virtue of the fact that the porous flame holder 106 is hot under normal, steady state operating conditions. As such, the body 202 of the porous flame holder 106 will generally not abstract sufficient heat from a flame travelling through a gas passage 110 to quench the flame even if the dimension D_D is below the classically defined quenching distance.

Since this is a thermal argument, actual flashback can occur through the quenching distance if the orifice is very hot—for example, if a premixed burner reservoir is receiving radiant heat from a hot furnace, e.g., a premix burner in ethylene service. But even so, in general the quenching distance does not change dramatically inasmuch as the flow of premixed fuel and air in the gaseous mixture 104 tend to cool the upstream end 206 of the walls 204 defining the gas passages 110. In one range of embodiments, the plurality of gas passages 110 each have a lateral dimension D_L between 0.05 inch and 1.0 inch. Preferably, the plurality of gas passages 110 have a lateral dimension D_L between 0.1 inch and 0.5 inch. For example the plurality of perforations can have a lateral dimension D_L of about 0.2 to 0.4 inch. In some embodiments, the gas passages 110 have substantially constant cross sectional areas. In other embodiments, the gas passages 110 can have non-constant cross sectional areas.

Moreover, the gaseous mixture 104 is typically introduced to the upstream surface 108 of the porous flame holder 106 at a dilution below the lean flammability limit of the fuel (more on that below).

The gas passages 110 each include a wall 204 configured to receive heat from an exothermic reacting portion of the gaseous mixture and from hot products of the exothermic reaction, transmit the heat toward an unreacted portion of the gaseous mixture, and output the heat adjacent to an unreacted portion of the gaseous mixture. The inventors have found that heat is apparently received from the combustion reaction along a heat receiving area 208 of the gas passage 110 walls 204 that starts at about $\frac{1}{3}$ of the length L from the upstream surface 108 to the downstream surface 112 of the porous flame holder 106 all the way to the downstream surface 112 of the perforated flame holder 106. Heat is conveyed upstream along the body 202 of the porous flame holder 106 through thermal conduction (diagrammatically depicted as 210), thermal radiation (diagrammatically depicted as 212), or (most likely) both thermal conduction 210 and thermal radiation 212. Net heat flow from the walls 204 of the gas passages 110 to the incoming gaseous mixture 104 apparently occurs in regions 206 that extend from the upstream surface 108 of the porous flame holder 106 to about $\frac{1}{3}$ of the length L from the upstream surface 108 to the downstream surface 112 of the porous flame holder.

The gas passages 110 are configured to fix a location of the combustion reaction in a flow of the gaseous mixture 104 by virtue of the heat transfer effects described above. At steady state operating conditions, the gas passages 110 of the

porous flame holder **106** are configured to hold the combustion reaction supported by the gaseous mixture **104** substantially between the upstream surface **108** and the downstream surface **112** of the porous flame holder **106**.

During start-up, after the porous flame holder **106** has been preheated, but before equilibrium is reached, the inventors have noted blue flames extending downstream of the downstream surface **112** of the porous flame holder **106**. Even in these conditions, however, it is believed that a majority, i.e. over 50%, of the combustion reaction occurs between the upstream surface **108** and the downstream surface **112** of the porous flame holder **106**.

In other experiments, the inventors found that a portion of the combustion reaction can occur in a region near and immediately upstream from the upstream surface **108** of the porous flame holder **106**. This is believed to be due to a combination of conduction (or convection) from the upstream ends of the porous flame holder body **202** defining the walls **204** of the gas passages and flow stagnation in the gaseous mixture **104** caused by aerodynamic effects of the upstream ends of the flame holder body **202**. Aside from transient effects, such as pulsing, the majority of the combustion reaction is carried within the porous flame holder **106**, between the upstream surface **108** and the downstream surface **112**.

The plurality of gas passages **110** can include a plurality of elongated squares. In another embodiment, the plurality of gas passages **110** can include a plurality of elongated hexagons. The porous flame holder **106** can be formed from VERSAGRID® ceramic honeycomb, available from Applied Ceramics, Inc. of Doraville, S.C., (illustrated in FIG. **1B**) for example. In another embodiment, the porous flame holder **106** can be formed from a plurality of tubes bundled together. In another embodiment, the porous flame holder **106** can be formed from an interleaved crimped surface and flat surface coiled together. In another embodiment, the porous flame holder **106** can be formed as a reticulated fiber mesh. In another embodiment, the porous flame holder **106** can be formed as a structured or unstructured packing. In another embodiment, the porous flame holder **106** can be formed as holes drilled in a fiber reinforced refractory material (illustrated in FIG. **1A**).

In some embodiments the gas passages **110** can pass from the upstream surface **108** of the porous flame holder **106** to the downstream surface **112** of the porous flame holder **106** separately. In other embodiments the gas passages can pass from the upstream surface **108** to the downstream surface **112** of the porous flame holder with intersections configured to allow comingling if the gaseous mixture **104** or the combustion reaction supported by the gaseous mixture within the porous flame holder **106**. The porous flame holder **106** can be constructed of a solid body or of a composite body, such as reticulated fibers or layered sheets.

The porous flame holder **106** can be formed from a refractory material such as alumina. Additionally or alternatively, the porous flame holder **106** can be formed from cordierite or mullite. In some embodiments, the porous flame holder **106** can be formed from a fiber reinforced cast refractory material. In another embodiment, the porous flame holder **106** can be formed from a metal superalloy such as Hastelloy or Inconel.

The inventors found that for a given flow velocity, a larger dimension D_L in a gas passage **110** requires a larger length L of the gas passage **110** to reach the lowest NOx production. For tested combinations, the length L was equal to the distance between the upstream surface **108** and downstream surface **112** of the porous flame holder **106**. Similarly,

smaller D_L was found to operate effectively with a smaller gas passage length L . Lengths L tested by the inventors range from about 1 inch to 8 inches, with the larger porous flame holder **106** thicknesses having larger dimension gas passages **110**. E.g., 2-inch thick porous flame holders **106** were tested at gas passage sizes corresponding to densities ranging from about 16 gas passages **110** per square inch (nominally a 0.25 inch square gas passage **110**) to 100 gas passages per square inch (nominally a 0.10 inch square gas passage **110**). A six inch thick porous flame holder **106** was tested at a gas passage density of 4 gas passages **110** per square inch (nominally a 0.5 inch square gas passage **110**) to 16 gas passages **110** per square inch.

The inventors tested porous flame holders **106** having void fractions between about 0.3 to greater than 0.7. The void fraction of a porous flame holder **106** is defined as the total area of all gas passages **110**, divided by a total area bounded by the upstream surface **108**. Some porous flame holder **106** embodiments can operate with less than 0.10 void fraction or more than 0.90 void fraction, but the inventors believe that a porous flame holder **106** having a void fraction between these limits is preferable. At a low void fraction, a porous flame holder **106** may exhibit undesirable pressure drop in the flowing gaseous mixture **104**. Between the two tested void fractions, the 0.70 void fraction porous flame holder **106**, illustrated in FIG. **1B**, produced lower NOx concentration than did the 0.30 void fraction porous flame holder **106** illustrated in FIG. **1A**.

The inventors have achieved stable heat outputs ranging from about 0.114 MBTU/(hr* ft^2) (million BTUs per hour per square foot of porous flame holder surface) to 1.2 MBTU/(hr* ft^2) while maintaining 3% O₂ in the stack. The inventors believe higher (stable) heat fluxes will probably be achieved with sufficient heat load to maintain a steady state temperature.

FIG. **3** is a graphically depiction **300** of representative emissions performance achieved by the burner of FIGS. **1A**, **1B**, and **2** in one experiment (but representative of many experiments). The inventors performed numerous observations and measurements of the illustrated effect. Data from a particular run is shown in FIG. **3**, wherein oxides of nitrogen (NOx) concentration **302** and carbon monoxide (CO) concentration **304** were monitored at the flue with 3% oxygen (O₂) stack concentration. The flue gas temperature was 1600° F.

During a 7-minute start-up period **306**, the porous flame holder was preheated to reach a start-up temperature characterized by between 800° F. and 1200° F. stack temperature. Visually, this corresponded to a bright reddish orange glow at the center of the porous flame holder **106** of the embodiment **101** shown in FIG. **1B**. Pre-heating was performed by a conventional flame held immediately adjacent to a fuel nozzle (depicted as **114a**, **114b**, **114** in FIGS. **1A** and **1B**). After 7 minutes, the start-up flame was blown off the start-up flame holder (one variant of a start-up flame holder **116** is depicted in FIG. **1A**), and a gaseous mixture (see **104**) of diluted fuel and combustion air transferred to the porous flame holder (see **106**) where, owing to the elevated temperature of the porous flame holder, the mixture immediately ignited.

Upon flame transfer, NOx concentration dropped precipitously from about 120 parts per million (ppm) characteristic of a conventional (start-up) flame to under 10 ppm. During a transition period **308** from 7-minutes to 10-minutes after start-up, NOx concentration asymptotically approached 5 ppm. After several minutes, NOx output reduced to 2 ppm or lower. During the transition period, fuel flow rate was

gradually increased to reach about 1 MBTU/(hr*ft²). During the transition period 308, the porous flame holder equilibrated to a steady state temperature distribution characterized by a bright orange glow. It is also shown that CO emissions were reduced to about 2 ppm.

Subsequent experiments were conducted wherein the NOx concentration was reduced below the 1 ppm detection limit of the flue gas sensor.

FIG. 4 is a diagram 400 of fuel and combustion air mixing used to produce a gaseous mixture 104, according to an embodiment. The diagram shows a fuel rich region 402 with an adjacent oxidant rich region 404. Another oxidant rich region (not shown) would be present to the left of the fuel rich region 402, but is omitted for simplicity. While the fuel and combustion air flow through a dilution region (see D_D in FIGS. 1A and 1B), mixing occurs in vortex cores 406 that may, to a first approximation, be regarded as being uniform in composition, having a fuel dilution between that of the fuel rich region 402 and the oxidant rich region 404. Between the vortex cores lies a Taylor layer 408, which at higher respective concentrations of fuel and oxidant than the concentrations in the vortex cores 406. As the stream 104 flows toward the porous flame holder (see 106), relatively pure combustion air and relatively pure fuel are engulfed by the vortex cores 406 in relatively "big gulps". As the stream 104 flows upward, more and more of the Taylor layer 408 is engulfed by the vortex cores to cause the maximum concentration of fuel and air, respectively, to diminish.

The first step in mixing is entrainment. In a free shear flow, such as the turbulent jet, ambient fluid 404 is entrained into the jet 402 by the large-scale engulfment of tongues of ambient fluid. The edges of the tongues are subsequently convoluted by the turbulence into progressively smaller-scale convolutions. According to the leading theory of turbulent mixing, the mixed fluid resides in only two places, the Taylor layer 408 associated with the strain rate of the largest eddies, and the Batchelor layer associated with the strain rate of the smallest eddies. The thicknesses of the Taylor and Batchelor layers are given by the plane, strained, laminar flame solution of Marble. From experiments in gaseous and aqueous shear layers, it turns out that the amount of mixed fluid in the Taylor layer 408 is about equal to that in the Batchelor layer in a gas flow. The fluid mixed at the Batchelor scale accumulates in the vortex cores and becomes essentially volume-filling, as sketched in FIG. 4.

Referring to FIGS. 1A and 1B, according to an embodiment, the fuel nozzle assembly 102 is configured to mix the gaseous mixture 104 sufficiently to substantially destroy Taylor layers 408 between pure fuel and air carrying the oxidant. In a non-premixed burner 100, 101 illustrated by FIGS. 1A and 1B, the fuel nozzle assembly 102 includes one or more fuel nozzles 114, 114a, 114b configured to output substantially pure fuel. The fuel nozzle assembly 102 further includes a combustion air source 118a, 118b, 118 configured to provide combustion air. In a non-premixed system, the combustion air may be provided as natural draft or by forced draft using, e.g., a blower. As the fuel from the fuel nozzles 114a, 114b, 114 travels upward through a dilution distance D_D , it entrains combustion air, which causes dilution. Typically the fuel jet expands at a 15° solid angle (shown diagrammatically in FIG. 1B), with the entirety of expansion corresponding to additional combustion air being incorporated into the fuel.

According to an embodiment, the fuel nozzle assembly 102 is configured to output the gaseous mixture 104 (including fuel and oxidant) to the porous flame holder 106 with sufficient air or flue gas to cause the gaseous mixture 104 to

be fuel-lean of a stoichiometric mixture. In some embodiments, the fuel nozzle assembly 102 is configured to output the gaseous mixture 104 substantially at a lean flammability limit of the fuel. The lean fuel and air mixture 104 can be used to reduce combustion temperature inside the porous flame holder 106.

The porous flame holder 106 has been found to output a significant amount of heat from the combustion reaction as thermal radiation 212. In other words, the porous flame holder 106 is configured to radiate heat away from the reacting portion of the gaseous mixture. In other words, the porous flame holder 106 is configured to radiate heat away from the combustion reaction supported by the gaseous mixture 104. The porous flame holder 106 can be configured to radiate heat away from the combustion reaction sufficiently to cause the gaseous mixture 104 to burn at or below 2000° F., wherein the gaseous mixture 104 consists essentially of air, methane, and flue gas from the combustion reaction. In some embodiments, the porous flame holder 106 is configured to radiate heat away from the combustion reaction sufficiently to cause the gaseous mixture 104 to burn at about 1700° F., wherein the gaseous mixture consists essentially of air, methane, and flue gas from the combustion reaction. Another aspect of the porous flame holder 106 is that it causes combustion to be completed in a very short time, which reduces the output of thermal NOx.

As described above, the burner 100, 101 can include a start-up flame holder 116 configured to hold the combustion reaction having a richer fuel mixture than the porous flame holder 106, wherein the start-up flame holder 116 is configured to support the combustion reaction at a location configured to pre-heat the porous flame holder 106.

The fuel nozzle assembly 102 can include a single fuel nozzle 114 or a plurality of nozzles 114a, 114b configured to output substantially pure fuel. The fuel nozzle assembly 102 can include an air source 118a, 118b, 118 configured to output substantially pure air. The fuel nozzle assembly 102 and porous flame holder 106 can be disposed to define a mixing zone D_D configured to allow mixing of the substantially pure fuel with the substantially pure air to produce a uniform gaseous mixture at the upstream surface 108 of the porous flame holder 106.

Alternatively, the burner 100, 101 can be configured as a pre-mix burner. In a pre-mix burner, the fuel nozzle assembly 102 includes a premixing chamber (not shown) operatively coupled to one or more fuel nozzles 114a, 114b, 114 and one or more air sources 118a, 118b, 118, and configured to uniformly mix fuel and air to form the gaseous mixture 104. In a pre-mix burner 100, 101, a flame arrestor can be disposed between the mixing chamber and the porous flame holder to prevent flashback.

Referring to FIG. 1A, a burner system 100 may include several spatial and temporal features surrounding a conventional nozzle 114a, 114b. The geometric elements can include a lower story tile 120, and the porous flame holder 106 supported above the lower story tile 120 by a support structure 122. When the fuel is directed to the lower story tile 120, the resulting recirculation of the hot gas anchors the combustion reaction in that region. This start-up mode is maintained until a characteristic temperature is attained. For example, the characteristic temperature could be the temperature of the porous flame holder 106 or some related temperature such as the ultimate system operation temperature or a switching temperature measured at the flue. Once the characteristic temperature is attained, a switching operation is initiated, which then anchors the combustion reaction at the porous flame holder 106. Combustion in the latter

configuration dramatically reduces NO_x via a variety of mechanisms. For example, so-called thermal NO_x is a function of three main factors: flame temperature, oxygen concentration, and time under such favorable NO_x formation conditions. By increasing the amount of entrained flue gas, oxygen concentration and NO_x-forming species are diluted. The greater entrainment also provides more thorough mixing and shorter flames and reduced time to form NO_x. Inasmuch as flame chemistry is much faster than fuel/air mixing, flame length in diffusion flames is primarily determined by the mixing of fuel and air to form a flammable mixture. By premixing the fuel and air before combustion on the upperstory tile, flame length is reduced. Since flame length is one parameter that limits the total heat output, its decrease allows for greater firing rate in a given furnace leading to greater process throughput.

As indicated above, the porous flame holder **106** plays two critical roles, according to embodiments. First, the porous flame holder **106** acts as a flame holder, fixing the ignition location. Second, the porous flame holder efficiently radiates energy away from the hot combustion products, thereby cooling them. The greater thermal conductivity of the porous flame holder **106** as compared to the flue gas also homogenizes flame temperature. Inasmuch as thermal NO_x formation is exponentially related to flame temperature, small volumes of higher temperature gas (so-called "hot spots") can contribute to the lion's share of NO_x formation. Therefore, a flame with fewer hot spots will generate less NO_x *ceteris paribus*.

Referring to FIG. **2**, the hot combustion products transfer heat to the walls **104** of the porous flame gas passages **110**. Each wall **104** may then radiates infrared and visible energy. Some of that energy is radiated upstream. Combined with thermal conduction **210** through the perforated flame holder **106**, thermal radiation **212** heats each gas passage wall in a region **206** and provides a heating source for the oncoming cold reactants flowing into the porous flame holder **106**. By transferring the energy upstream, the residence time of the unburned reactants in the heated zone is increased, which may improve the turndown ratio of the burner.

Thermal radiation **212** also is output toward the furnace walls and/or other heat loads in the system. As a consequence, the temperature of the combustion products is promptly reduced, lowering the NO_x emissions. The solid surface of the perforated flame holder **106** is a much better radiator than the combustion products, especially if there is little soot.

Unlike a diffusion flame, the burner **100**, **101** of FIGS. **1A**, **1B**, and **2** delays combustion until sufficient entrainment of air and flue gas insures low combustion reaction temperatures and low NO_x emissions. During operation, combustion occurs in the burner far downstream from the fuel nozzle(s) **114a**, **114b**, **114**, so that every parcel of injected fuel has mixed with sufficient air to be lean of stoichiometric. The Taylor layers (see FIG. **4**, **408**) between pure fuel and air have been destroyed, and the vortex cores **406** are lean. Consequently, the temperature of the combustion products is relatively low.

FIG. **5** is a flowchart showing a method **500** of operating the burners of FIGS. **1A**, **1B**, and **2** to lower combustion NO_x, according to an embodiment. Prior to the steps of the method **500**, a start-up temperature and/or operating temperature is established in the porous flame holder. As described above, establishing a start-up temperature can be performed by holding a conventional flame held by a start-up flame holder in a position to cause heating of the porous flame holder.

At step **502**, fuel and oxidant are combined into a gaseous mixture. The gaseous mixture may include a turbulently-moving gas. In one embodiment, step **502** includes outputting a substantially pure fuel jet from a fuel nozzle and outputting substantially pure combustion air from a combustion air source. The fuel jet can entrain the air (including oxygen as the oxidant). The air may be introduced by natural convection through an air source concentric to the fuel nozzle, for example. In other embodiments, the air is introduced by forced convection from a blower.

Outputting a substantially pure fuel jet can include operating a fuel nozzle assembly including a plurality of nozzles each configured to output substantially pure fuel. The fuel nozzle assembly can include an air source configured to output substantially pure air.

Proceeding to step **504**, time is allowed for the gaseous mixture to evolve to a better-mixed state. Step **504** may include time for turbulent mixing, for example. In the embodiments of FIGS. **1A** and **1B**, the mixing time is provided by the dilution distance D_D through which the gaseous mixture travels. The fuel nozzle assembly and placement of the porous flame holder can define a mixing zone configured to allow the time for mixing of substantially pure fuel with substantially pure air to produce a uniform gaseous mix at the porous flame holder.

Step **504** may include allowing time sufficient to substantially destroy Taylor layers between pure fuel and air carrying the oxidant. Step **504** may include providing sufficient air or flue gas to cause the fuel to be at a lean of stoichiometric mixture. Step **504** can include forming the mixture to be substantially at a lean flammability limit of the fuel.

In step **506** the gaseous mixture is introduced into a porous flame holder. Step **506** can include simultaneously introducing the gaseous mixture to an upstream surface of the porous flame holder such that the gaseous mixture enters a plurality of gas passages simultaneously. The porous flame holder can be formed from a refractory material such as alumina, mullite, and/or cordierite. In another embodiment, the porous flame holder can be formed from a superalloy such as Hastelloy or Inconel.

In step **508**, the gaseous mixture is combusted inside a porous flame holder. The porous flame holder (see **106**) is configured to conduct heat sufficiently to cause combustion to be supported inside the porous flame holder. The step of combusting the mixture includes passing the mixture (and a combustion reaction, and combustion reaction products) through gas passages of the porous flame holder. In an embodiment, the gas passages have a minimum dimension equal to or greater than a quenching distance associated with the fuel. The gas passages may have substantially constant cross sectional areas. In an embodiment, the gas passages are formed at a pitch of 10 per lineal inch across an upstream surface of the porous flame holder or at a lower pitch (larger cells). For example, the gas passages can be formed at a pitch of 4 per inch or higher across two dimensions of the porous flame holder.

Step **508** includes receiving heat evolved from the combustion reaction through walls of the gas passages. The heat can be radiated, conducted, or radiated and conducted toward the upstream surface of the porous flame holder. The hot walls then transfer heat toward unreacted portions of the gaseous mixture to heat the gaseous mixture up to a combustion temperature. Step **508** also includes radiating heat away from the porous flame holder to cool the combustion sufficient to cause the mixture to burn at or below 2000° F. In some embodiments, the porous flame holder radiates

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sufficient heat away from the combustion reaction to cause the mixture to burn at or below 1700° F.

Optionally, steps 502 and 504 can include operating a fuel pre-mixer. Operating the fuel pre-mixer can include using a pre-mixing chamber to uniformly mix the fuel and oxidant mixture and disposing a flame arrestor between the mixing chamber and the porous flame holder.

While various aspects and embodiments have been disclosed herein, other aspects and embodiments are contemplated. The various aspects and embodiments disclosed herein are for purposes of illustration and are not intended to be limiting, with the true scope and spirit being indicated by the following claims.

What is claimed is:

1. A burner, comprising:

a fuel nozzle assembly configured to output at least one jet of substantially pure fuel configured to entrain oxidant to provide a gaseous mixture including the fuel and the oxidant; and

a porous flame holder defining gas passages each having a minimum lateral dimension equal to or greater than a fuel quenching distance, the gas passages being aligned to receive the gaseous mixture from the fuel nozzle assembly, the gas passages being configured to pass a combustion reaction supported by the gaseous mixture; wherein the gas passages each include a respective wall arranged and configured to receive heat from a reacting portion of the gaseous mixture at a heat receiving area of the wall, convey the received heat via at least one of thermal conduction and thermal radiation to an upstream portion of the wall adjacent to an unreacted portion of the gaseous mixture, and transfer the heat to the unreacted portion of the gaseous mixture within the gas passage; and

wherein the fuel nozzle assembly is configured to mix the gaseous mixture sufficiently to, as the gaseous mixture approaches the porous flame holder, substantially destroy Taylor layers between the substantially pure fuel and air carrying the oxidant in a structurally uninterrupted space between the fuel nozzle assembly and the porous flame holder.

2. The burner of claim 1, wherein the gas passages are further configured to fix a location of the combustion reaction in a flow of the gaseous mixture.

3. The burner of claim 1, wherein the fuel nozzle assembly is configured to output the gaseous mixture to the porous flame holder with sufficient air or flue gas to cause the fuel to be at a lean of stoichiometric mixture.

4. The burner of claim 3, wherein the fuel nozzle assembly is configured to output the gaseous mixture substantially at a lean flammability limit of the fuel.

5. The burner of claim 1, wherein the porous flame holder is configured to radiate heat away from the combustion reaction supported by the gaseous mixture.

6. The burner of claim 1, wherein the porous flame holder is configured to radiate heat away from the combustion reaction sufficiently to cause the gaseous mixture to burn at or below 2000° F., wherein the gaseous mixture consists essentially of air, methane, and flue gas from the combustion reaction.

7. The burner of claim 6, wherein the porous flame holder is configured to radiate heat away from the combustion reaction sufficiently to cause the gaseous mixture to burn at about 1700° F., wherein the gaseous mixture consists essentially of air, methane, and flue gas from the combustion reaction.

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8. The burner of claim 1, wherein the gas passages have substantially constant cross-sectional areas.

9. The burner of claim 1, wherein the gas passages are formed at a pitch of 10 per inch or less across two dimensions of the porous flame holder.

10. A burner, comprising:

a fuel nozzle assembly configured to output at least one jet of substantially pure fuel configured to entrain oxidant to provide a gaseous mixture including the fuel and the oxidant; and

a porous flame holder defining gas passages each having a minimum lateral dimension equal to or greater than a fuel quenching distance, the gas passages being aligned to receive the gaseous mixture from the fuel nozzle assembly, the gas passages being configured to pass a combustion reaction supported by the gaseous mixture; wherein the gas passages each include respective walls arranged and configured to receive heat from a reacting portion of the gaseous mixture, radiate or conduct the heat toward an upstream portion of the wall adjacent to an unreacted portion of the gaseous mixture, and transfer the heat to the unreacted portion of the gaseous mixture within the gas passage;

wherein the fuel nozzle assembly is configured to, as the gaseous mixture approaches the porous flame holder, mix the gaseous mixture sufficiently to substantially destroy Taylor layers between the substantially pure fuel and air carrying the oxidant in a structurally uninterrupted space between the fuel nozzle assembly and the porous flame holder; and

wherein the gas passages are formed at a pitch of 4 per inch or more than 4 per inch across two dimensions of the porous flame holder.

11. The burner of claim 1, wherein the porous flame holder is formed from a refractory material.

12. The burner of claim 1, wherein the fuel nozzle assembly includes a plurality of nozzles configured to output substantially pure fuel.

13. The burner of claim 12,

wherein the fuel nozzle assembly includes an air source configured to output substantially pure air; and wherein the fuel nozzle assembly defines a mixing zone configured to allow mixing of the substantially pure fuel with the substantially pure air to produce a uniform gaseous mixture at the porous flame holder.

14. A method of lowering combustion NO_x, comprising: outputting, from a fuel nozzle assembly, at least one jet of substantially pure fuel and entrained oxidant toward a porous flame holder; then

allowing time for a mixture of the fuel and oxidant to evolve to a better-mixed state while the fuel and the oxidant traverse a structurally uninterrupted dilution distance between the fuel nozzle assembly and the porous flame holder, the dilution distance selected to permit said allowing of time; and then

combusting the mixture inside the porous flame holder; wherein walls of gas passages of the porous flame holder are each configured to radiate or conduct heat, received from the combusting of the mixture at a heat receiving region of the respective wall, to an upstream region of the wall adjacent to an unreacted portion of the mixture, and transfer the heat to the unreacted portion of the mixture sufficiently to cause the combustion to be supported inside the porous flame holder, and

wherein the dilution distance between the fuel nozzle assembly and the porous flame holder is selected to permit substantial destruction of Taylor layers between

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the substantially pure fuel and air carrying the oxidant as the mixture reaches the porous flame holder.

15. The method of claim 14, wherein the step of combusting the mixture comprises passing the mixture through the gas passages of the porous flame holder, the gas passages having a minimum lateral dimension equal to or greater than a fuel quenching distance of the mixture.

16. The method of claim 15, wherein the gas passages have substantially constant cross-sectional areas.

17. The method of claim 15, wherein the gas passages are formed at a pitch of 10 per inch or less across two dimensions of the porous flame holder.

18. A method of lowering combustion NO_x, comprising: outputting, from a fuel nozzle assembly, at least one jet of substantially pure fuel and entrained oxidant toward a porous flame holder; then

allowing time for a mixture of the fuel and the oxidant to evolve to a better-mixed state while the fuel and the oxidant traverse a structurally uninterrupted dilution distance between the fuel nozzle assembly and the porous flame holder, the dilution distance selected to permit said allowing of time; and then

combusting the mixture inside the porous flame holder; wherein walls of gas passages of the porous flame holder are each configured to radiate or conduct heat, received from the combusting of the mixture, to a portion of the mixture in another portion of the gas passage sufficiently to cause the combustion to be supported inside the porous flame holder,

wherein the dilution distance between the fuel nozzle assembly and the porous flame holder is selected to permit substantial destruction of Taylor layers between the substantially pure fuel and air carrying the oxidant as the mixture reaches the porous flame holder,

wherein the step of combusting the mixture comprises passing the mixture through the gas passages of the porous flame holder, the gas passages having a minimum lateral dimension equal to or greater than a fuel quenching distance of the mixture; and

wherein the gas passages are formed at a pitch of 4 per inch or more across two dimensions of the porous flame holder.

19. The method of claim 15, wherein the gas passages include a wall configured to receive heat from a reacting portion of the mixture, radiate the heat toward an unreacted portion of the mixture, and receive the radiated heat adjacent to the unreacted portion of the mixture.

20. The burner of claim 14, wherein the step of outputting fuel and oxidant further comprises providing sufficient air or flue gas to cause the fuel to be at a lean of stoichiometric mixture.

21. The burner of claim 20, further comprising forming the better-mixed state of the mixture to be substantially at a lean flammability limit of the fuel.

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22. The method of claim 14, further comprising configuring a radiative property of the porous flame holder sufficient to radiate heat away from the combustion to cause the mixture to burn at or below 2000° F., when the gaseous mixture consists essentially of air, methane, and flue gas from a combustion reaction.

23. The method of claim 22, further comprising configuring the radiative property of the porous flame holder sufficient to radiate the heat away from the combustion to cause the mixture to burn at or below 1700° F., when the gaseous mixture consists essentially of the air, the methane, and the flue gas from the combustion reaction.

24. The method of claim 14, wherein the fuel nozzle assembly in the step of outputting fuel and oxidant includes a plurality of nozzles configured to output substantially pure fuel.

25. The method of claim 24,

wherein the fuel nozzle assembly includes an air source configured to output substantially pure air; and

wherein the fuel nozzle assembly defines a mixing zone configured to allow mixing of the substantially pure fuel with the substantially pure air to produce a uniform gaseous mix at the porous flame holder.

26. The burner of claim 14, wherein the porous flame holder is formed from a refractory material.

27. A diluted fuel burner, comprising:

a fuel nozzle assembly configured to output at least one jet of substantially pure fuel and to entrain oxidant to produce a gaseous mixture including the fuel and the oxidant;

a porous flame holder defining gas passages aligned to receive the gaseous mixture from the fuel nozzle assembly, the gas passages being configured to carry a combustion reaction supported by the gaseous mixture; and

a porous flame holder support structure configured to hold the porous flame holder away from the fuel nozzle assembly at a structurally uninterrupted dilution distance selected to allow dilution of the gaseous mixture; wherein the dilution distance is selected to permit substantial destruction of Taylor layers between the substantially pure fuel and air carrying the oxidant as the mixture approaches the porous flame holder.

28. The diluted fuel burner of claim 27, wherein the gas passages include respective walls each configured to receive heat from a reacting portion of the gaseous mixture, radiate or conduct the heat toward an upstream region of the wall adjacent to an unreacted portion of the gaseous mixture, and transfer the heat into the unreacted portion of the gaseous mixture within the gas passage.

29. The burner of claim 1, wherein the gas passages have non-constant cross-sectional areas.

30. The burner of claim 1, further comprising a natural draft, non-pressurized oxidant source.

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